(FILE 'HOME' ENTERED AT 20:04:27 ON 23 SEP 2004)

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, AQUALINE, ANABSTR, ANTE, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS, DDFB, DDFU, DGENE, ...' ENTERED AT 20:07:02 ON 23 SEP 2004 SEA (THIOESTERAS?(S)OBES?) OR (HYDROLAS?(S)COA?(S)OBES?)

4 FILE BIOSIS

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- FILE BIOTECHDS
- 4 FILE BIOTECHNO
- 5 FILE CABA
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- 3 FILE LIFESCI
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- 9 FILE SCISEARCH
- 1 FILE TOXCENTER
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- 1 FILE USPAT2
- 2 FILE WPIDS
- 2 FILE WPINDEX
- 1 FILE NLDB QUE (THIOESTERAS?(S) OBES?) OR (HYDROLAS?(S) COA?(S) OBES?)

LI

- FILE 'DGENE, EMBASE, CAPLUS, USPATFULL, SCISEARCH, ESBIOBASE, CABA,
- BIOSIS, BIOTECHNO, MEDLINE' ENTERED AT 20:09:32 ON 23 SEP 2004 108 S (THIOESTERAS?(S)OBES?) OR (HYDROLAS?(S)COA?(S)OBES?) L2
- L3 76 DÙP REM L2 (32 DÙPLICATÉS RÈMOVED)
- L4 7 S L3 (S)ADIPOS?

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        SEP 01
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         SEP 01
NEWS 14 SEP 14 STN Patent Forum to be held October 13, 2004, in Iselin, NJ
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FULL ESTIMATED COST

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- 74 FILES SEARCHED...

=> d rank

F1

- 1 FILE NLDB
- 26 FILES HAVE ONE OR MORE ANSWERS, . 77 FILES SEARCHED IN STNINDEX
- QUE (THIOESTERAS?(S) OBES?) OR (HYDROLAS?(S) COA?(S) OBES?) L1
- 42 **DGENE** F2 13 EMBASE F3 10 CAPLUS F4 10 USPATFULL 9 SCISEARCH 7 F6 **ESBIOBASE** F7 5 CABA F8 4 BIOSIS F9 4 BIOTECHNO F10 4 MEDLINE F11 **BIOTECHABS** F12 3 BIOTECHDS F13 3 DDFU F14 3 DRUGU F15 3 GENBANK F16 3 LIFESCI F17 3 PASCAL F18 3* FEDRIP F19 2 IFIPAT F20 2 PHIN F21 WPIDS F22 2 WPINDEX F23 **EMBAL**

F24 1 TOXCENTER F25 1 USPAT2 F26 1 NLDB

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=> dup rem 12 DUPLICATE IS NOT AVAILABLE IN 'DGENE'. ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE PROCESSING COMPLETED FOR L2 76 DUP REM L2 (32 DUPLICATES REMOVED)

=> d ti 13 1-76

L3ANSWER 1 OF 76 USPATFULL on STN TΙ Novel proteins and nucleic acids encoding same

ANSWER 2 OF 76 USPATFULL on STN L3TI Gene expression in bladder tumors

ANSWER 3 OF 76 USPATFULL on STN L3

Methods of diagnosis of breast cancer, compositions and methods of ΤI screening for modulators of breast cancer

1.3 ANSWER 4 OF 76 USPATFULL on STN ΤI

Inhibition of fatty acid synthase by beta-lactones and other compounds for inhibition of cellular proliferation

L3 ANSWER 5 OF 76 USPATFULL on STN

ТT Tumor necrosis factor receptor 2

ANSWER 6 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. 1.3 on STN DUPLICATE 1

- TI A fatty acid synthase blockade induces tumor cell-cycle arrest by down-regulating Skp2.
- L3 ANSWER 7 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. ON STN DUPLICATE 2
- TI Orlistat Is a Novel Inhibitor of Fatty Acid Synthase with Antitumor Activity.
- L3 ANSWER 8 OF 76 USPATFULL on STN
- TI Protein-protein interactions in adipocyte cells (3)
- L3 ANSWER 9 OF 76 USPATFULL on STN
- TI BFIT compositions and methods of use
- L3 ANSWER 10 OF 76 USPATFULL on STN
- TI Human Transcriptomes
- L3 ANSWER 11 OF 76 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on STN
- TI Regulation of human S-acyl fatty acid synthase thioesterase-like enzyme.
- L3 ANSWER 12 OF 76 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation. on STN DUPLICATE 3
- TI Human uncoupling protein-3 and obesity: An update
- L3 ANSWER 13 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Hepatic protein expression of lean mice and obese diabetic mice treated with peroxisome proliferator-activated receptor activators. [Erratum to document cited in CA139:95192]
- L3 ANSWER 14 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Gene expression profiles of nondiabetic and diabetic obese mice suggest a role of hepatic lipogenic capacity in diabetes susceptibility
- L3 ANSWER 15 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Hepatic protein expression of lean mice and obese diabetic mice treated with peroxisome proliferator-activated receptor activators
- L3 ANSWER 16 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
- TI cDNAs encoding human and mouse brown fat inducible acyl CoA thioesterases and their use in treatment of metabolic disorders
- L3 ANSWER 17 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Cloning, sequencing and regulation of human S-Acyl fatty acid synthase thioesterase-like protein
- L3 ANSWER 18 OF 76 USPATFULL on STN
- TI 56939, a novel human acyl-CoA thioesterase family member and uses thereof
- L3 ANSWER 19 OF 76 USPATFULL on STN
- TI Regulation of human S-acyl fatty acid synthase thioesterase-like enzyme
- L3 ANSWER 20 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. ON STN DUPLICATE 4
- TI Glucose-dependent regulation of cholesterol ester metabolism in macrophages by insulin and leptin.
- L3 ANSWER 21 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Microarray profiling of skeletal muscle tissues from equally obese, non-diabetic insulin-sensitive and insulin-resistant Pima Indians
- L3 ANSWER 22 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. on STN DUPLICATE 5
- TI De novo expression of uncoupling protein 3 is associated to enhanced mitochondrial thioesterase-1 expression and fatty acid metabolism in liver of fenofibrate-treated rats.
- L3 ANSWER 23 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.

 On STN DUPLICATE 6
- TI BFIT, a unique acyl-CoA thioesterase induced in thermogenic

brown adipose tissue: Cloning, organization of the human gene and assessment of a potential link to **obesity**.

- L3 ANSWER 24 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.
 ON STN DUPLICATE 7
- TI Structure and function of ASP, the human homolog of the mouse agouti gene.
- L3 ANSWER 25 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. ON STN DUPLICATE 8
- TI Effect of dietary n-3 and n-6 polyunsaturated fatty acids on lipidmetabolizing enzymes in obese rat liver.
- L3 ANSWER 26 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Subcellular fractionation evidence for a putative peroxisome-mitochondrion attachment in the liver of normal and genetically obese (ob/ob and db/db) mice
- L3 ANSWER 27 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. ON STN DUPLICATE 9
- TI Metabolic effects of coconut, safflower, or menhaden oil feeding in lean and obese Zucker rats.
- L3 ANSWER 28 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. ON STN DUPLICATE 10
- TI Comparison of dehydroepiandrosterone and clofibric acid treatments in obese Zucker rats.
- L3 ANSWER 29 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Effect of short-term DHEA administration on liver metabolism of lean and obese rats.
- L3 ANSWER 30 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN DUPLICATE 11
- TI Effect of short-term DHEA administration on liver metabolism of lean and obese rats
- L3 ANSWER 31 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.
- TI A comparison of the effects of dehydroepiandrosterone treatment to ad libitum and pair-feeding in the obese Zucker rat.
- L3 ANSWER 32 OF 76 CABA COPYRIGHT 2004 CABI on STN
- TI A comparison of the effects of dehydroepiandrosterone treatment to ad libitum and pair-feeding in the obese Zucker rat.
- L3 ANSWER 33 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. ON STN DUPLICATE 12
- TI Metabolic consequences of dehydroepiandrosterone in lean and obese adult Zucker rats.
- L3 ANSWER 34 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. on STN
- TI Lipid synthesis in lactating mammary gland.
- L3 ANSWER 35 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible **thioesterase** (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, **obesity** or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 36 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT ON STN
- TI New isolated of brown fat inducible **thioesterase** (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, **obesity** or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 37 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible **thioesterase** (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, **obesity** or cachexia associated with HIV infection, sepsis, trauma or cancer.

- ANSWER 38 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New isolated human acyl-CoA thioesterase polypeptide 56939, useful for treating cardiovascular, liver, metabolic, brain, kidney, and cellular proliferative and/or differentiative disorders -
- ANSWER 39 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New purified human S-acyl fatty acid synthase thioesterase-like
 enzyme, useful for identifying modulators of enzyme activity for treating
 cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 40 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New purified human S-acyl fatty acid synthase thioesterase-like
 enzyme, useful for identifying modulators of enzyme activity for treating
 cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 41 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New purified human S-acyl fatty acid synthase thioesterase-like
 enzyme, useful for identifying modulators of enzyme activity for treating
 cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 42 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New purified human S-acyl fatty acid synthase thioesterase-like
 enzyme, useful for identifying modulators of enzyme activity for treating
 cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 43 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New purified human S-acyl fatty acid synthase thioesterase-like
 enzyme, useful for identifying modulators of enzyme activity for treating
 cardiovascular disease, diabetes, obesity and hyperlipidaemia
- L3 ANSWER 44 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
 TI New isolated of brown fat inducible thioesterase (BFIT)
 polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with
 HIV infection, sepsis, trauma or cancer.
- ANSWER 45 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New isolated of brown fat inducible thioesterase (BFIT)
 polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- ANSWER 46 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New isolated of brown fat inducible thioesterase (BFIT)
 polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- ANSWER 47 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT ON STN

 New isolated of brown fat inducible thioesterase (BFIT)
 polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- ANSWER 48 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

 New isolated of brown fat inducible thioesterase (BFIT)
 polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- ANSWER 49 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT ON STN

 New isolated of brown fat inducible thioesterase (BFIT)
 polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.

- L3 ANSWER 50 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible **thioesterase** (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, **obesity** or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 51 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 52 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 53 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- New isolated of brown fat inducible thioesterase (BFIT)
 polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with
 HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 54 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 55 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible thioesterase (BFIT)
 polypeptides, useful for diagnosing and treating patients with metabolic
 disease, e.g. diabetes, obesity or cachexia associated with
 HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 56 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 57 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- New isolated of brown fat inducible **thioesterase** (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, **obesity** or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 58 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 59 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 60 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection, sepsis, trauma or cancer.
- L3 ANSWER 61 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- TI New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with

HIV infection, sepsis, trauma or cancer.

ANSWER 62 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN L3 TΤ New isolated of brown fat inducible thioesterase (BFIT) polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, **obesity** or cachexia associated with HIV infection, sepsis, trauma or cancer.

ANSWER 63 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

New isolated of brown fat inducible thioesterase (BFIT) TТ polypeptides, useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, **obesity** or cachexia associated with HIV infection, sepsis, trauma or cancer.

- L3 ANSWER 64 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
- New isolated human acyl-CoA thioesterase polypeptide 56939, useful for ΤI treating cardiovascular, liver, metabolic, brain, kidney, and cellular proliferative and/or differentiative disorders
- L3 ANSWER 65 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT ON STN New purified human S-acyl fatty acid synthase thioesterase-like enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia
- L3 ANSWER 66 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN New purified human S-acyl fatty acid synthase thioesterase-like TI

enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia

- ANSWER 67 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN TI New purified human S-acyl fatty acid synthase thioesterase-like
- enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia
- L_3 ANSWER 68 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN New purified human S-acyl fatty acid synthase thioesterase-like TI enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 69 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN 1.3 New purified human S-acyl fatty acid synthase thioesterase-like TΙ enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 70 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN L3 New purified human S-acyl fatty acid synthase thioesterase-like TT enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 71 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT ON STN L3TTNew purified human S-acyl fatty acid synthase thioesterase-like enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 72 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN L3New purified human S-acyl fatty acid synthase thioesterase-like enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia
- ANSWER 73 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN L3New purified human S-acyl fatty acid synthase thioesterase-like TΙ enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia

- ANSWER 74 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT On STN L3
- ΤI New purified human S-acyl fatty acid synthase thioesterase-like

enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia

ANSWER 75 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN L3

New purified human S-acyl fatty acid synthase thioesterase-like TT enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia

ANSWER 76 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN L3

ΤI New purified human S-acyl fatty acid synthase thioesterase-like enzyme, useful for identifying modulators of enzyme activity for treating cardiovascular disease, diabetes, obesity and hyperlipidaemia

=> d 13 9 11-12 16-17 18 23 62 76 ibib abs

ANSWER 9 OF 76 USPATFULL on STN 1.3

ACCESSION NUMBER:

2003:312634 USPATFULL

TITLE:

BFIT compositions and methods of use

INVENTOR(S):

Adams, Sean H., Randolph Township, NJ, UNITED STATES Chui, Clarissa J., San Francisco, CA, UNITED STATES Goddard, Audrey D., San Francisco, CA, UNITED STATES Grimaldi, J. Christopher, San Francisco, CA, UNITED

STATES

Lewin, David A., New Haven, CT, UNITED STATES

KIND NUMBER DATE US 2003220238 20031127 A1 US 2002-55624 A1 20020122 (10)

APPLICATION INFO .:

NUMBER DATE

PRIORITY INFORMATION:

PATENT INFORMATION:

US 2001-263362P 20010122 (60)

DOCUMENT TYPE:

Utility

FILE SEGMENT:

APPLICATION

LEGAL REPRESENTATIVE:

SONNENSCHEIN NATH & ROSENTHAL LLP, P.O. BOX 061080,

WACKER DRIVE STATION, SEARS TOWER, CHICAGO, IL,

60606-1080

NUMBER OF CLAIMS:

31 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS:

7 Drawing Page(s)

LINE COUNT: 2751

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Isolated polynucleotides encoding brown fat inducible thioesterase (BFIT) polypeptides and the polypeptides are provided. Methods of using these polynucleotides and polypeptides are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 11 OF 76 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on STN

ACCESSION NUMBER:

2003:377866 BIOSIS

DOCUMENT NUMBER:

PREV200300377866

TITLE:

Regulation of human S-acyl fatty acid synthase

thioesterase-like enzyme.

AUTHOR(S):

Xiao, Yonghong [Inventor, Reprint Author]; Towler, Eric Marshall [Inventor]; Eveleigh, Jamie Frederick [Inventor]

CORPORATE SOURCE: Cambridge, MA, USA

ASSIGNEE: Bayer Aktiengesellschaft, Leverkusen, Germany

PATENT INFORMATION: US 6593099 July 15, 2003

SOURCE:

Official Gazette of the United States Patent and Trademark

Office Patents, (July 15 2003) Vol. 1272, No. 3. http://www.uspto.gov/web/menu/patdata.html. e-file. ISSN: 0098-1133 (ISSN print).

DOCUMENT TYPE: LANGUAGE:

Patent English

ENTRY DATE:

Entered STN: 13 Aug 2003

Last Updated on STN: 13 Aug 2003

Reagents which regulate human S-acyl fatty acid synthase thioesterase-like enzyme and reagents which bind to human S-acyl fatty acid synthase thioesterase-like enzyme gene products can play a role in preventing, ameliorating, or correcting dysfunctions or diseases including, but not limited to cardiovascular disease, hyperlipidemia, obesity, and diabetes.

ANSWER 12 OF 76 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation. L_3 DUPLICATE 3 on STN

ACCESSION NUMBER:

2004:62300 SCISEARCH

THE GENUINE ARTICLE: 759FF

AUTHOR:

Human uncoupling protein-3 and obesity: An update Hesselink M K C (Reprint); Mensink M; Schrauwen P

CORPORATE SOURCE:

Maastricht Univ, Dept Movement Sci, POB 616, NL-6200 MD Maastricht, Netherlands (Reprint); Maastricht Univ, Dept

Movement Sci, NL-6200 MD Maastricht, Netherlands;

Maastricht Univ, Dept Human Biol, Nutr & Toxicol Res Inst

Maastricht, NL-6200 MD Maastricht, Netherlands

COUNTRY OF AUTHOR:

SOURCE:

OBESITY RESEARCH, (DEC 2003) Vol. 11, No. 12, pp.

Netherlands 1429-1443.

Publisher: NORTH AMER ASSOC STUDY OBESITY, 8630 FENTON ST,

SUITE 918, SILVER SPRING, MD 20910 USA.

ISSN: 1071-7323.

DOCUMENT TYPE:

General Review; Journal

LANGUAGE:

English

REFERENCE COUNT: 94

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS The cloning of the uncoupling protein (UCP)I homologs UCP2 and UCP3 has AB raised considerable interest in the mechanism. The expression of UCP3 mainly in skeletal muscle mitochondria and the potency of the skeletal muscle as a thermogenic organ made UCP3 an attractive target for studies toward manipulation of energy expenditure to fight disorders such as obesity and type 2 diabetes. Overexpressing UCP3 in mice resulted in lean, hyperphagic mice. However, the lack of an apparent phenotype in mice lacking UCP3 triggered the search for alternative functions of UCP3. The observation that fatty acid levels significantly affect UCP3 expression has given UCP3 a position in fatty acid handling and/or oxidation. Emerging data indicate that the primary physiological role of UCP3 may be the mitochondrial handling of fatty acids rather than the regulation of energy expenditure through thermogenesis. It has been proposed that UCP3 functions to export fatty acid anions away from the mitochondrial matrix. In doing so, fatty acids are exchanged with protons, explaining the uncoupling activity of UCP3. The exported fatty acid anions may originate from hydrolysis of fatty acid esters by a mitochondrial thioesterase, or they may have entered the mitochondria as nonesterified fatty acids by incorporating into and flip-flopping across the mitochondrial inner membrane. Regardless of the origin of the fatty acid anions, this putative function of UCP3 might be of great importance in protecting mitochondria against fatty acid accumulation and may help to maintain muscular fat oxidative capacity.

ANSWER 16 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:676171 CAPLUS

DOCUMENT NUMBER:

137:212048

TITLE:

SOURCE:

LANGUAGE:

cDNAs encoding human and mouse brown fat inducible

acyl CoA thioesterases and their use in treatment of

metabolic disorders

INVENTOR(S):

Adams, Sean H.; Goddard, Audrey D.; Grimaldi, J.

Christopher; Chui, Clarissa J.

PATENT ASSIGNEE(S):

Curagen Corporation, USA; Genentech, Inc.

PCT Int. Appl., 92 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent English

FAMILY ACC. NUM. COUNT: 1

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PATENT NO.
                         KIND
                                DATE
                                            APPLICATION NO.
                                                                    DATE
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     WO 2002068619
                          A2
                                20020906
                                            WO 2002-US1783
                                                                    20020122
     WO 2002068619
                          A3
                                20030710
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             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
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     US 2003220238
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                          A1
                                            US 2002-55624
                                                                    20020122
     EP 1368462
                          A2
                                20031210
                                            EP 2002-705889
                                                                   20020122
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
         R:
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
PRIORITY APPLN. INFO.:
                                            US 2001-263362P
                                                                P 20010122
                                            WO 2002-US1783
                                                                W 20020122
     Isolated polynucleotides encoding brown fat inducible thioesterase (BFIT)
     polypeptides and the polypeptides are provided. BFIT mRNA species was
     induced 2-3-fold above the control following cold exposure (4.degree.),
     and repressed .apprxeq. 70% by warm acclimation (33.degree.C, 3 wk)
     compared with controls (22.degree.C). BFIT was mapped to syntenic regions
     of chromosomes 1 (human) and 4 (mouse) assocd. with body fatness and
     diet-induced obesity, potentially linking a deficit of BFIT activity with
     exacerbation of these traits. Consistent with this notion, BFIT mRNA was
     significantly higher (\sim 1.6-2-fold) in the brown adipose tissue of
     obesity-resistant compared with obesity-prone mice fed a high-fat diet,
     and was 2.5-fold higher in controls compared with ob/ob mice. The strong,
     cold-inducible brown adipose tissue expression in mice suggests that BFIT
     supports the transition of this tissue towards increased metabolic
     activity, probably through alteration of intracellular fatty acyl-CoA
     concn. Methods of using these polynucleotides and polypeptides are also
     provided.
    ANSWER 17 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:
                         2002:10659 CAPLUS
DOCUMENT NUMBER:
                         136:80920
TITLE:
                         Cloning, sequencing and regulation of human S-Acyl
                         fatty acid synthase thioesterase-like protein
                         Xiao, Yonghong
INVENTOR(S):
PATENT ASSIGNEE(S):
                         Bayer Aktiengesellschaft, Germany
SOURCE:
                         PCT Int. Appl., 123 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
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English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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	20020008				2002		1	WO 2	001-	 EP72	 97		2	0010	 626
WO 2	20020008				2002										
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	ВJ,	CF, C	G, CI,	CM,	GA.	GN.	GW.	ML.	MR.	NE.	SN.	TD,	TG,	110,	DI,
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AB Reagents which regulate human S-acyl fatty acid synthase thioesterase-like enzyme and reagents which bind to human S-acyl fatty acid synthase thioesterase-like enzyme gene products can play a role in preventing, ameliorating, or correcting dysfunctions or diseases including, but not limited to cardiovascular disease, hyperlipidemia, obesity, and diabetes.

L3 ANSWER 18 OF 76 USPATFULL on STN

ACCESSION NUMBER: 2002:294282 USPATFULL

TITLE: 56939, a novel human acyl-CoA thioesterase family

member and uses thereof

INVENTOR(S): Meyers, Rachel A., Newton, MA, UNITED STATES

NUMBER DATE

PRIORITY INFORMATION: US 2000-220040P 20000721 (60)

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: LOUIS MYERS, FISH & RICHARDSON P.C., 225 Franklin

Street, Boston, MA, 02110-2804

NUMBER OF CLAIMS: 19 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 2 Drawing Page(s)

LINE COUNT: 5096

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention provides isolated nucleic acids molecules, designated 56939 nucleic acid molecules, which encode novel acyl-CoA thioesterase family members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 56939 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 56939 gene has been introduced or disrupted. The invention still further provides isolated 56939 proteins, fusion proteins, antigenic peptides and anti-56939 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 23 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED. On STN DUPLICATE 6

ACCESSION NUMBER:

2001412436 EMBASE

TITLE:

BFIT, a unique acyl-CoA thioesterase induced in

thermogenic brown adipose tissue: Cloning, organization of the human gene and assessment of a potential link to

obesity.

AUTHOR:

SOURCE:

Adams S.H.; Chui C.; Schilbach S.L.; Yu X.X.; Goddard A.D.;

Grimaldi J.C.; Lee J.; Dowd P.; Colman S.; Lewin D.A.

CORPORATE SOURCE:

S.H. Adams, Metabolic Disease Pharmacology Dept., Novartis Pharmaceuticals Corporation, 556 Morris Avenue, Summit, NJ

07901, United States. sean.adams@pharma.novartis.com Biochemical Journal, (15 Nov 2001) 360/1 (135-142).

Refs: 44

ISSN: 0264-6021 CODEN: BIJOAK

COUNTRY:
DOCUMENT TYPE:
FILE SEGMENT:

United Kingdom

Journal; Article

005 General Pathology and Pathological Anatomy

029 Clinical Biochemistry

LANGUAGE: English SUMMARY LANGUAGE: English

We hypothesized that certain proteins encoded by temperature-responsive genes in brown adipose tissue (BAT) contribute to the remarkable metabolic shifts observed in this tissue, thus prompting a differential mRNA expression analysis to identify candidates involved in this process in mouse BAT. An mRNA species corresponding to a novel partial-length gene was found to be induced 2-3-fold above the control following cold exposure (4.degree.C), and repressed .apprxeq. 70 % by warm acclimation (33.degree.C, 3 weeks) compared with controls (22.degree.C). The gene displayed robust BAT expression (i.e. .apprxeq. 7-100-fold higher than

other tissues in controls). The full-length murine gene encodes a 594 amino acid (.apprxeq. 67 kDa) open reading frame with significant homology to the human hypothetical acyl-CoA thioesterase KIAA0707. Based on cold-inducibility of the gene and the presence of two acyl-CoA thioesterase domains, we termed the protein brown-fat-inducible thioesterase (BFIT). Subsequent analyses and cloning efforts revealed the presence of a novel splice variant in humans (termed hBFIT2), encoding the orthologue to the murine BAT gene. BFIT was mapped to syntenic regions of chromosomes 1 (human) and 4 (mouse) associated with body fatness and diet-induced obesity, potentially linking a deficit of BFIT activity with exacerbation of these traits. Consistent with this notion, BFIT mRNA was significantly higher (.apprxeq. 1.6-2-fold) in the BAT of obesity-resistant compared with obesity-prone mice fed a high-fat diet, and was 2.5-fold higher in controls compared with ob/ob mice. Its strong, cold-inducible BAT expression in mice suggests that BFIT supports the transition of this tissue towards increased metabolic activity, probably through alteration of intracellular fatty acyl-CoA concentration.

L3 ANSWER 62 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT ON STN

ACCESSION NUMBER: ADH13925 DNA DGENE

TITLE: New isolated of brown fat inducible thioesterase

(BFIT) polypeptides, useful for diagnosing and treating

patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with HIV infection,

sepsis, trauma or cancer.

INVENTOR: Adams S H; Goddard A D; Grimaldi J C; Chui C J

PATENT ASSIGNEE: (CURA-N) CURAGEN CORP.

(GETH) GENENTECH INC.

PATENT INFO: WO 2002068619 A2 20020906 92p

APPLICATION INFO: WO 2002-US1783 20020122 PRIORITY INFO: US 2001-263362P 20010122

DOCUMENT TYPE: Patent LANGUAGE: English

OTHER SOURCE: 2002-698669 [75]

DESCRIPTION: Murine brown fat inducible thioesterase PCR primer rev SEQ ID

NO:17.

AN ADH13925 DNA DGENE

The invention relates to a novel isolated brown fat inducible thioesterase (BFIT) polypeptide. A polypeptide of the invention has anorectic, antidiabetic, and immunomodulator activity. A polynucleotide of the invention may have a use in gene therapy. The BFIT polynucleotides and polypeptides are useful for diagnosing and treating patients with metabolic disease, e.g. diabetes, obesity or cachexia associated with human immunodeficiency virus (HIV) infection, sepsis, trauma or cancer. The antibodies and methods are useful for screening agonists or antagonists that regulate or affect BFIT polynucleotides and polypeptides. The present sequence is used in the exemplification of the invention.

L3 ANSWER 76 OF 76 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

ACCESSION NUMBER: ABK13701 cDNA DGENE

TITLE: New purified human S-acyl fatty acid synthase

thioesterase-like enzyme, useful for identifying

modulators of enzyme activity for treating cardiovascular

disease, diabetes, obesity and hyperlipidaemia

INVENTOR: Xiao Y

PATENT ASSIGNEE: (FARB) BAYER AG.

PATENT INFO: WO 2002000855 A2 20020103 123p

APPLICATION INFO: WO 2001-EP7297 20010626
PRIORITY INFO: US 2000-214012P 20000626
US 2000-255148P 20001214

DOCUMENT TYPE: Patent LANGUAGE: English

OTHER SOURCE: 2002-130886 [17] CROSS REFERENCES: P-PSDB: AAU75501

DESCRIPTION: DNA encoding s-acyl fatty acid synthase thioesterase-like

enzyme #1.

AN ABK13701 cDNA DGENE

AB The invention describes a purified human S-acyl fatty acid synthase thioesterase (SFST)-like enzyme (I). (I) is useful for screening

for agents which decrease or regulate the activity of an SFST-like enzyme and agents which decrease extracellular matrix degradation. The polynucleotide (II) is useful for detecting a polynucleotide which encodes (I) in a biological sample by formation of a hybridisation complex. A reagent modulating the activity or SFST-like enzyme or an antibody can also be used to detect the polypeptide or polynucleotide in a biological sample. A reagent modulating the activity of (I) or (II) is also useful for treating a SFST-like enzyme dysfunction related disease condition such as cardiovascular disease, hyperlipidaemia, obesity, anorexia, cachexia, wasting disorders, appetite suppression, appetite enhancement, bulimia or diabetes. The cardiovascular diseases treated by the above mentioned methods and reagents include congestive heart failure, myocardial infarction, ischaemic diseases of the heart, atrial and ventricular arrhythmia, hypertensive vascular diseases, and peripheral vascular diseases. (I) is useful in diagnostic assays for detecting diseases and abnormalities related to presence of mutations in the nucleic acid sequences which encode the enzyme. This sequence encodes a human s-acyl fatty acid synthase thioesterase-like enzyme, described in the method of the invention.

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L1

L2 L3 (FILE 'HOME' ENTERED AT 20:04:27 ON 23 SEP 2004)

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, AQUALINE, ANABSTR, ANTE, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS, DDFB, DDFU, DGENE, ...' ENTERED AT 20:07:02 ON 23 SEP 2004 SEA (THIOESTERAS?(S)OBES?) OR (HYDROLAS?(S)COA?(S)OBES?)

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     FILE CABA
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     FILE CAPLUS
     FILE DDFU
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42
     FILE DGENE
     FILE DRUGU
1
     FILE EMBAL
13
     FILE EMBASE
     FILE ESBIOBASE
    FILE FEDRIP
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     FILE GENBANK
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     FILE IFIPAT
 3
     FILE LIFESCI
     FILE MEDLINE
    FILE PASCAL
3
     FILE PHIN
9
     FILE SCISEARCH
1
     FILE TOXCENTER
10
     FILE USPATFULL
    FILE USPAT2
1
     FILE WPIDS
     FILE WPINDEX
     FILE NLDB
 QUE (THIOESTERAS?(S) OBES?) OR (HYDROLAS?(S) COA?(S) OBES?)
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FILE 'DGENE, EMBASE, CAPLUS, USPATFULL, SCISEARCH, ESBIOBASE, CABA, BIOSIS, BIOTECHNO, MEDLINE' ENTERED AT 20:09:32 ON 23 SEP 2004

108 S (THIOESTERAS?(S)OBES?) OR (HYDROLAS?(S)COA?(S)OBES?)

76 DUP REM L2 (32 DUPLICATES REMOVED)

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FIELD CODE - 'AND' OPERATOR ASSUMED 'L24 (S)ADIPOS?'

PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH

FIELD CODE - 'AND' OPERATOR ASSUMED 'L30 (S)ADIPOS?'

PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH
FIELD CODE - 'AND' OPERATOR ASSUMED 'L32 (S)ADIPOS?'
L4 7 L3 (S) ADIPOS?

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L1

(FILE 'HOME' ENTERED AT 20:04:27 ON 23 SEP 2004)

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, AQUALINE, ANABSTR, ANTE, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS, DDFB, DDFU, DGENE, ...' ENTERED AT 20:07:02 ON 23 SEP 2004 SEA (THIOESTERAS?(S)OBES?) OR (HYDROLAS?(S)COA?(S)OBES?)

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    FILE WPINDEX
    FILE NLDB
 QUE (THIOESTERAS?(S) OBES?) OR (HYDROLAS?(S) COA?(S) OBES?)
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FILE 'DGENE, EMBASE, CAPLUS, USPATFULL, SCISEARCH, ESBIOBASE, CABA, BIOSIS, BIOTECHNO, MEDLINE' ENTERED AT 20:09:32 ON 23 SEP 2004

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L3 76 DUP REM L2 (32 DUPLICATES REMOVED)

L4 7 S L3 (S)ADIPOS?

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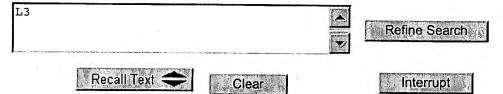
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US Patents Full-Text Database
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JPO Abstracts Database
Derwent World Patents Index
IBM Technical Disclosure Bulletins

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Search History

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Search Results - Record(s) 1 through 3 of 3 returned.

☐ 1. Document ID: US 20040043382 A1

Using default format because multiple data bases are involved.

L2: Entry 1 of 3

File: PGPB

Mar 4, 2004

PGPUB-DOCUMENT-NUMBER: 20040043382

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040043382 A1

TITLE: Novel proteins and nucleic acids encoding same

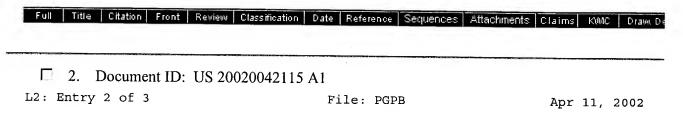
PUBLICATION-DATE: March 4, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Padigaru, Muralidhara	Branford	CT	US	
Spytek, Kimberly A.	New Haven	CT	US	
Shenoy, Suresh G.	Branford	CT	US	
Taupier, Raymond J. JR.	East Haven	CT	US	
Pena, Carol E. A.	New Haven	CT	US	
Li, Li	Branford	CT	US	
Zerhusen, Bryan D.	Branford	CT	US	
Gusev, Vladimir Y.	Madison	CT	US	
Ji, Weizhen	Branford	CT	US	
Gorman, Linda	Branford	CT	US	
Miller, Charles E.	Guilford	CT	US	
Kekuda, Ramesh	Norwalk .	CT	US	
Patturajan, Meera	Branford	CT	US	
Gangolli, Esha A.	Madison	CT	US	
Vernet, Corine A.M.	Branford	CT	US	
Guo, Xiaojia Sasha	Branford	CT	US	
Tchernev, Velizar T.	Branford	CT	US	
Fernandes, Elma R.	Branford	CT	US	
Casman, Stacie J.	North Haven	CT	US	
Malyankar, Uriel M.	Branford	CT	US ·	
Gerlach, Valerie	Branford	CT	US	
Liu, Yi	San Diego	CA	US	
Anderson, David W.	Branford	CT	US	
Spaderna, Steven K.	Berlin	CT	US	
Catterton, Elina	Madison	CT	US	
Leite, Mario W.	Milford	CT	US	

Zhong, Haihong	Guilford	СТ	US
Alsobrook, John P. II	Madison	CT	US
Lepley, Denise M.	Branford	CT	US
Rieger, Daniel K.	Branford	CT	US
Burgess, Catherine E.	Wethersfield	CT	US

US-CL-CURRENT: $\underline{435}/\underline{6}$; $\underline{435}/\underline{183}$, $\underline{435}/\underline{320.1}$, $\underline{435}/\underline{325}$, $\underline{435}/\underline{69.1}$, $\underline{530}/\underline{350}$, $\underline{536}/\underline{23.2}$



PGPUB-DOCUMENT-NUMBER: 20020042115

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020042115 A1

TITLE: Regulation of human S-acyl fatty acid synthase thioesterase-like enzyme

PUBLICATION-DATE: April 11, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Xiao, Yonghong

Cambridge

MA

US

US-CL-CURRENT: 435/183; 435/320.1, 435/325, 435/69.1, 536/23.2

Full	Title Cita	ation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWC	Draw, De
	3. Doc	umer	nt ID:	US 65	93099 B2				a)			
L2: Er	ntry 3 d	of 3				\mathbf{F}^{\sharp}	ile: USP	T		Jul	15,	2003

US-PAT-NO: 6593099

DOCUMENT-IDENTIFIER: US 6593099 B2

TITLE: Regulation of human S-acyl fatty acid synthase thioesterase-like enzyme

DATE-ISSUED: July 15, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Xiao; Yonghong Cambridge MA
Towler; Eric Marshall New Haven CT
Eveleigh; Jamie Frederick West Haven CT

US-CL-CURRENT: 435/19; 435/18, 435/197, 536/23.2

Full	Title	Citation	Front	Review	Classification	Date	Reference	Scopulline	Attachine	de Claims	KWIC	Draw, D
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